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(54) BIODEGRADABLE SCLERA PLUG

BIOLOGISCH ABBAUBARER SKLERASTOPFEN

BOUCHON DE SCLEROTIQUE BIODEGRADABLE

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Description

[0001] The present invention relates to a biodegradable medical device to deliver a medication to a specific internal site, which is made of lactic acid polymer and contains a drug. Moreover, the present invention relates to a medical device that can be used to treat or prevent diseases of the retina or to promote a recovery from damage after vitreous surgery while taking advantage of the sustained release of a drug.

[0002] A vitreous body is one of intraocular tissues that is viscous, transparent, and gel-like. A vitreous body consists mainly of water and includes components comprising collagen fibers and hyaluronic acid. The retina is the most inner tissue layer of the eye. It covers a vitreous body and is in direct contact therewith. Diseases of the retina are generally intractable and some of the diseases are accompanied by opacity of the vitreous body. Examples of such diseases are retinal hemorrhage caused by various damage, proliferative vitreoretinopathy accompanied by vascularization and proliferation of retinal cells, retinal detachment and retinoblastoma.

[0003] Eye diseases are generally treated by instillation of drugs. However, drugs can barely reach the retina or vitreous body by instillation. Moreover, the blood-aqueous barrier reduces delivery of effective concentration of a drug to the retina or a vitreous body even by a systemic administration, e.g., intravenous administration. It is not practical to inject a drug directly into a vitreous body, because injection of a high concentration of a drug may cause disorders of intraocular tissues and the procedure for repeated injection is laborious and may cause infection.

[0004] Some diseases are treated by intraocular surgery. For example, in the case of vitreous hemorrhage, the opaque vitreous body is excised, and in the case of proliferative vitreoretinopathy, the proliferative tissues and the vitreous body are excised. In the case of retinal detachment, the vitreous body is excised and an artificial vitreous body such as silicone is used to hold the retina steady. In vitreous surgery, the sclera that is the eyeball wall is incised at three sites; a site to irrigate physiological irrigating solution into the eye, a site to insert a guide that is used to light up the intraocular portion, and a site to insert tools for excision of a vitreous body, such as a cutter. A scleral plug made of metal may be used temporarily to insert tools into the eye or remove them (Am. J. Ophthalmol. 91, 797 (1981)), but the incisions are finally sutured to complete surgery.

[0005] For medicinal treatment of diseases of the retina or diseases of the vitreous body, a direct injection of drugs into the vitreous body is used. Methods for treating diseases have been studied in which microspheres or liposomes were used to release drugs slowly (Inv. Ophthalmol. Vis. Sci. 32, 1785 (1991)).

[0006] Recently, polymers from lactic acid have been studied as biodegradable polymers. Application

studies of such polymers are reported, for example, as a substance to fix fractures of bones (Japanese Unexamined Patent Publication 29663/91), as a suture for surgery (Japanese Unexamined Patent Publication 501109/92), or as a vehicle for an intraocular implantation (Japanese Unexamined Patent Publication 22516/88). Sustained release of drugs with the use of poly(lactic acid) has also been studied (J. Med. Chem. 16, 897 (1973)). Furthermore, sustained release preparations for intraocular implantation using low molecular-weight poly(lactic acid) have been reported (Japanese Unexamined Patent Publication 17370/93).

[0007] US-patent 4,863,457 discloses an implant for delivering continuous controlled supply for medication to a specific internal site of an eye. This medical device consists of a biodegradable material impregnated with therapeutic agents, which are released and can be diffused into the sites to be treated as the implant erodes.

[0008] Specifically the implant known from the US-patent comprises a more or less flat, circular base portion to be positioned in the sclera below the conjunctival membrane located beneath eyelids, and a tapered stem portion formed as a hook or an anchor to assist in retaining the implant in place.

[0009] However, despite the provision of a hook or anchor the polymer material of the implant may possibly be moved inside the intraocular tissues, which might cause a bad influence on surrounding tissues and may also effect a visual field and contact tissues of lens to cause cataract depending on the site where the polymer is implanted. Accordingly, additionally to the provision of the hook or anchor the known implant needs a fixation to the sclera with a suture or the like. However, such sutures are a burden for the patient and accordingly to be avoided.

[0010] Accordingly, it is the object of the invention to provide a medical device of the above-mentioned kind, by which drugs may be released gradually to the intraocular tissues for a long period of time avoiding the need for an additional fixing by suture or the like.

[0011] According to the present invention this object is solved in that the device is a nail-like scleral plug and has a head portion and a shaft portion, wherein the shaft portion only of the scleral plug is insertable into the vitreous body through the sclera such that the head portion remains outside the vitreous body, whereby the shaft portion is absorbable into the vitreous body so that the drug is released gradually into the vitreous body from the scleral plug.

[0012] The present invention makes use of the fact, that the diseases of the retina often treated by intraocular surgery in which a sclera has incision sites. Namely, according to the present invention the scleral plug is fixed so as not to move without any specific means, because it is inserted into the decision site of the sclera and firmly fits therein by making use of the shrinkage force of the sclera.

[0013] Further, as the result of our research on material for a scleral plug that need not be removed after surgery, we found that by using a lactic acid copolymer as the plug material, a scleral plug that is gradually degraded in the eye tissues and absorbed into the tissues, is safe to the highly sensitive tissues of the eye, can be prepared. Meanwhile, various properties are further desired to be provided to a scleral plug. Desired properties include maintaining the function to plug a scleral incisions for a required period, being degraded and absorbed in the required period, being easily handled during surgery, and so on. First, we examined a range of molecular weights for the lactic acid copolymer and the molar ratio of lactic acid units in the lactic acid copolymer which should satisfy the desired conditions, and found that it is preferable that the weight-average molecular weight of the lactic acid copolymer is in the range from 10,000 to 1,000,000 and the molar ratio of lactic acid units in the lactic acid copolymer is in the range from 50 to 100 mole %. Further, as a result of our study on the shape of a scleral plug, we found that it is preferable for a scleral plug to have a nail-like shape comprising a head portion, which prevents the plug from dropping into the inner site of eye, and a shaft portion, which is inserted through a scleral incision into the vitreous body. In particular, it is preferable that the end of the shaft portion is pointed, i.e., it is an acute-angled shape, such as pyramidal or conical, to prevent complication damage which may be caused when the plug is inserted.

[0014] We also studied a method to release a drug gradually into a vitreous body, and found that the object to release a drug gradually can be attained by allowing the drug to be contained in a scleral plug made of lactic acid copolymer, inserting the plug through a scleral incision into the vitreous body, and putting the end portion of the plug into the vitreous body. The insertion of the plug is usually carried out at a site of pars plana. It is possible to design the scleral plug of the present invention to enable a drug to be released depending on a desired releasing period by altering the molecular weight and/or the molar ratio of lactic acid units of the lactic acid copolymer. Because the plug is made of a biodegradable polymer, we found that the scleral plug of the present invention need not be removed after the drug is completely released, which could make it possible to attain effectively the object of the present invention. We also found that the scleral plug of the present invention could be inserted not only through an incision from vitreous surgery but also through a small opening directly made into a vitreous body through the sclera because the scleral plug of this invention can easily be inserted into the vitreous body through the sclera. Therefore, the scleral plug of the present invention can be used to treat or prevent diseases of the retina even if vitreous surgery is not carried out. If a very long drug-releasing period is needed, additional scleral plugs of the present invention can be inserted. If a high concen-

tration of the drug is needed, a plurality of the plugs of the present invention can be inserted simultaneously.

[0015] The present invention relates to a scleral plug made of lactic acid copolymer; a scleral plug containing a drug that is designed to gradually release the drug into a vitreous body; and a scleral plug that is used to treat or prevent diseases of the retina or to promote recovery from damage after vitreous surgery taking advantage of its sustained release effect for drug delivery. As used herein, "a lactic acid copolymer" generally means a copolymer comprising lactic acid units and glycolic acid units. However, malic acid, glyceric acid, or tartaric acid, etc. can also be used instead of glycolic acid. "A lactic acid copolymer" also includes a copolymer consisting of lactic acid units in a molar ratio of 100%, i.e., poly (lactic acid). A lactic acid unit may be in a L-, D-, or DL-form.

[0016] A scleral plug needs to be strong enough not to break or chip by manipulation with a pincette during surgery. Moreover, a scleral plug needs to have properties to release a drug slowly during the desired period of time for treatment, etc., and to be degraded in eye tissues and absorbed into the tissues afterwards. The properties of a scleral plug can be determined by the weight-average molecular weight of the copolymer, the ratio of lactic acid units and glycolic acid units (unless otherwise stated the ratio is represented on a molar basis), and the like. To simplify the following explanation hereinafter, a copolymer consisting of lactic acid units and glycolic acid units is taken as a sample of copolymers. The degree of degradation of the copolymer essentially depends upon the degree of crystallinity and water absorptivity of the copolymer. As either the ratio of lactic acid units or that of glycolic acid units becomes higher, the degree of crystallinity becomes higher and the degradation rate becomes lower. When the ratio of lactic acid units and glycolic acid units is 50/50, the degradation rate is the highest. As the ratio of glycolic acid units in the copolymer becomes higher, the characteristics of polyglycolic acid appear and result in an increase of hydrophilicity, because polyglycolic acid is slightly more hydrophilic than poly(lactic acid). Therefore, the water absorptivity becomes higher and the degradation rate becomes higher. If the ratio of glycolic acid units is over 50%, the plug is difficult to prepare, because the copolymer is not easily dissolved in the usual organic solvents that are used to prepare the plug. Therefore, in consideration of the above mentioned characteristics of glycolic acid units, the ratio of glycolic acid units is preferably 50% or less.

[0017] The degradation rate of the plug is generally in proportion to the weight-average molecular weight of the copolymer. Therefore, as the molecular weight becomes higher, the degradation rate becomes lower.

[0018] When the plug is prepared, the ratio of lactic acid units and glycolic acid units and the molecular weight of the copolymer can be selected by taking account of the above-mentioned characteristics of the

composition and functions required of the plug. The molar content of the lactic acid units in the copolymer for the scleral plug of the present invention is preferably 50 to 100%. The molar content of the glycolic acid units is preferably 0 to 50%. The molecular weight of the copolymer affects the strength of the plug. As the molecular weight becomes higher, the strength of the plug is increased. Taking into consideration the strength required for the plug, the molecular weight of the copolymer is preferably 10,000 or more. When the molecular weight is too high, the degradation rate of the plug becomes low and the molding of the plug becomes difficult. Accordingly the molecular weight of the copolymer is preferably 1,000,000 or less. Thus, the weight-average molecular weight of the copolymer is preferably from 10,000 to 1,000,000 and can be selected from the range depending on the application purposes.

[0019] The weight-average molecular weight of the copolymer and ratio of lactic acid units and glycolic acid units are determined depending upon the period of time to maintain the effective concentration of the drug. The period of time to maintain the effective concentration of the drug, i.e., a required releasing period, is mainly determined depending upon the disease to be treated, the symptoms, and the effects of the drug. The scleral plug of the present invention can be preferably used to treat various kinds of retina diseases or to promote recovery from damage after vitreous surgery. For example, proliferative vitreoretinopathy, viral infection, post-operative inflammation, postoperative infection can be treated or prevented. The releasing period of a drug can be controlled in the range from one week to six months by selecting the molecular weight of the copolymer or the ratio of lactic acid units and glycolic acid units.

[0020] If the desired releasing period is relatively short, such as one to two weeks, the molecular weight of the copolymer can be about 10,000 to 100,000 and preferably about 10,000 to 50,000, and more preferably 20,000 to 40,000. If the desired releasing period is about two weeks to one month, the molecular weight of the copolymer can be about 10,000 to 200,000 and preferably about 20,000 to 100,000, and more preferably 20,000 to 50,000. If the desired releasing period is long, i.e., more than one month, for example, one to six months, the molecular weight of the copolymer can be about 10,000 to 1,000,000 and preferably about 20,000 to 400,000, and more preferably 40,000 to 200,000. As described above, the releasing period of a drug can also be controlled by altering the molar ratio of lactic acid units and glycolic acid units. Taking into consideration the releasing period of the drug and the degree of degradation of the plug, an appropriate molecular weight and the ratio of lactic acid units and glycolic acid units in the copolymer are selected. The ratio of lactic acid units and glycolic acid units is in the range from 100/0 to 50/50.

[0021] In more detail, when the drug is doxorubicin hydrochloride, which is useful for proliferative vitre-

oretinopathy, the releasing period for the drug is preferably two weeks to one month. The molecular weight of the copolymer of the scleral plug is about 10,000 to 200,000, preferably about 20,000 to 100,000, more preferably about 20,000 to 50,000 and the ratio of lactic acid units and glycolic acid units is 100/0 to 50/50, preferably 80/20 to 50/50, more preferably about 80/20 to 70/30. When the drug is ganciclovir, which is useful for viral infection, the releasing period of the drug is more than one month, for example one to six months, preferably about four to six months. The molecular weight of the copolymer of the scleral plug is about 10,000 to 1,000,000, preferably about 20,000 to 400,000, more preferably about 40,000 to 200,000 and the molar ratio of lactic acid units and glycolic acid units is 100/0 to 50/50, preferably 80/20 to 50/50, more preferably about 80/20 to 70/30.

[0022] A plurality of the scleral plugs of the present invention can be used simultaneously and successively or additionally. Therefore, if a high concentration of the drug is needed for clinical treatment, a plurality of the plugs can be used simultaneously, and if a releasing period of the drug should be extended, the plugs can be used successively or additionally. Thus, even if a desired amount of the drug can be hardly contained in a piece of plug, a desired amount of the drug can be released into the vitreous body by using plugs simultaneously or successively.

[0023] Japanese Unexamined Patent Publication 17370/93 discloses the use of poly(lactic acid) in a sustained release preparation that can be implanted in eyes, but the poly(lactic acid) is low in average molecular weight (up to about 7000). Such low molecular-weight poly(lactic acid) can not be used as a material for the scleral plug, because the above-discussed desired properties cannot be obtained.

[0024] A preferred shape of the scleral plug is a nail-like shape comprising a head portion, which prevents the plug from dropping into the eye ball, and a shaft portion, which is inserted into a scleral incision. In particular, it is preferable that the end of the shaft portion be pointed, i.e., it is an acute-angled shape, such as, pyramidal or conical to prevent disease complication, which may be caused when the plug is inserted. Preferably, the head portion is formed in a hemispheric, disk-like or polygonal (e.g., hexagonal) shape and the shaft portion is formed in prismatic, columnar, or the like, shape. The following information on the sizes of the plug is given by way of explanation to make understanding the shape of the plug of the present invention easier. The length of the plug is normally about 6 mm, the diameter of the head portion is about 2 mm, the diameter or width of the shaft portion is about 1 mm, and the weight of the plug is about 9 mg. Of course, these sizes can be altered depending upon the amount of the drug to be contained. A release of the drug into the vitreous body can be attained by diffusion of the drug accompanying hydrolysis of the copolymer of the plug. The plug

is not required to be removed because it is degraded in eye tissues and absorbed into the tissues. The amount of the released drug can be adjusted to a level that can maintain the effective concentration in the vitreous body and does not cause retinal damage, since the retina is easily damaged by contact with a high concentration of drugs. The amount of the released drug can be controlled by selecting the content of the drug in the plug, the molecular weight of the copolymer and a ratio of lactic acid units and glycolic acid units. The amount of the drug to be contained in the plug can be selected depending on the period over which the effective concentration of the drug should be maintained. The period can be selected depending on the disease to be treated or prevented, the symptoms, the properties or effects of the drug, etc. The period to maintain the effective concentration of a drug is normally in the range of a week to six months. For example, when antitumor agents, such as doxorubicin hydrochloride, which can be used to treat proliferative vitreoretinopathy, are used, a standard releasing period of the drug is in the range from about two weeks to one month. When antiviral agents such as ganciclovir, which can be used to treat viral infections, are used, a standard releasing period of the drug is more than one month, e.g., about one month to six months. When antiinflammatory agents, such as steroids, which can be used to treat postoperative inflammation, are used, a standard releasing period of the drug is in the range from about two weeks to one month. When antibiotics, which can be used to treat postoperative infections, are used, a standard releasing period of the drug is in the range from about one week to two weeks. When antifungal agents, which can be used to treat mycotic infections, are used, a standard releasing period of the drug is more than one month, for example, in the range from about one month to two months. A variety of drugs that are useful to treat or prevent various diseases of the retina or to promote recovery from damage after vitreous surgery, can be used. The characteristic of the present invention is a technology to apply biodegradable lactic acid copolymers and the drug itself does not limit applications of the present invention. Various kinds of drugs such as antitumor agents, antibiotics, antiinflammatory agents, antiviral agents or antifungal agents can be used.

[0025] General methods of preparing the plug of the present invention comprises the steps of synthesizing a lactic acid copolymer according to a known method, dissolving the copolymer in an organic solvent, such as methylene chloride, acetonitrile, acetic acid, or the like, adding a drug to the mixture, removing the solvent, and forming a plug from the obtained powder. The lactic acid copolymer can be prepared by e.g., polymerizing lactic acid or the cyclic dimer thereof and a comonomer such as glycolic acid or the cyclic dimer thereof in the presence of a catalyst such as tin octylate and lauryl alcohol. A bulk polymerization is preferable, but the polymerization methods are not limited.

BRIEF DESCRIPTION OF THE DRAWINGS

[0026]

Figure 1 is a perspective view of the scleral plug prepared in the Productional example 1 below; and Figure 2 is a perspective view of the scleral plug prepared in the Productional example 2 below.

BEST MODE OF CARRYING OUT THE INVENTION

[0027] Processes for producing scleral plugs for sustained release of drugs, i.e., doxorubicin hydrochloride, which can be used to treat or prevent proliferative vitreoretinopathy, and ganciclovir, which can be used to treat or prevent viral infection, are exemplified below. The following examples are given to make the present invention clearly understood and shall not limit the scope of the present invention.

(Productional example 1)

[0028] First, an L-lactic acid/glycolic acid (in a molar ratio of 75/25) copolymer was synthesized in the presence of a catalyst consisting of tin octylate and lauryl alcohol by a bulk ring-opening polymerization. The weight-average molecular weight of the obtained copolymer was 38,000. After the copolymer (198 mg) was dissolved in 2 ml methylene chloride, doxorubicin hydrochloride (2 mg) was dispersed in the solution while stirring. The resulting dispersion was cast on a Teflon sheet to obtain an L-lactic acid/glycolic acid copolymer sheet containing doxorubicin hydrochloride. Then the scleral plug shown in Fig. 1 was formed from the sheet by a cutting and skiving process.

[0029] The scleral plug 4 is a nail-like shape consisting of a disk-like shaped head portion 1 and a shaft portion 3 of a square pillar. The end 2 of the shaft portion 3 is pyramidal.

(Productional example 2)

[0030] 1000mg of a copolymer (molar ratio of L-lactic acid unit/glycolic acid unit: 75/25, the weight-average molecular weight was 40,000) prepared as described in Productional example 1 and doxorubicin hydrochloride (5 mg) were dissolved in 5 ml of acetonitrile/water (9/1). The solvent was then removed from the solution by lyophilization. The resulting powder was formed into a scleral plug under heating to obtain the scleral plug shown in Fig. 2.

[0031] The scleral plug 14 is a nail-like shape consisting of a hemispheric head portion 11 and a columnar shaft portion 13. The end 12 of the shaft portion 13 is conical.

[0032] By similar methods to those as described above, a scleral plug which consists of L-lactic acid/glycolic acid in a molar ratio of 50/50, 70/30, or 80/20 and

having a weight-average molecular weight of 10,000, 20,000, 50,000, 70,000, 100,000, or 200,000; and which contains doxorubicin hydrochloride by 0.1%, 0.3%, 0.5%, 1%, or 2% can be obtained. By using D-lactic acid or DL-lactic acid instead of L-lactic acid, a similar scleral plug can be obtained.

(Productional example 3)

[0033] 900mg of a copolymer (DL-lactic acid/glycolic acid: 75/25, the weight-average molecular weight was 122,000) prepared as described in Productional example 1 and ganciclovir (100 mg) were dissolved in 15 ml of acetic acid and the mixture was lyophilized. The resulting powder was formed into a scleral plug under heating.

[0034] By similar methods to those described above, a scleral plug which consists of DL-lactic acid/glycolic acid in a molar ratio of 50/50, 70/30, or 80/20 and having a weight-average molecular weight of 10,000, 20,000, 40,000, 200,000, 400,000, or 1,000,000; and which contains ganciclovir by 1%, 2%, 5%, 10%, 15%, or 20% can be obtained. By using L-lactic acid or D-lactic acid instead of DL-lactic acid, a similar scleral plug can be obtained.

(Productional example 4)

[0035] 900mg of poly(L-lactic acid) (the weight-average molecular weight was 95,000) and ganciclovir (100 mg) were dissolved in 15 ml of acetic acid and then the mixture was lyophilized. The resulting powder was formed into a scleral plug under heating.

[0036] By similar methods to those described above, a scleral plug having a weight-average molecular weight of 10,000, 20,000, 40,000, or 200,000; and which contains ganciclovir by 1%, 2%, 5%, 10%, 15% or 20% can be obtained. By using poly (DL-lactic acid) or poly(D-lactic acid) instead of poly (L-lactic acid), a similar scleral plug can be obtained.

[0037] Sustained release effect of the drug, safety and degradation rate of the scleral plug which contains doxorubicin hydrochloride as an example of a drug were tested. The scleral plug containing doxorubicin hydrochloride prepared in Productional example 1 above was used.

(1) In vitro tests

[0038] The scleral plug was incubated in an isotonic phosphate buffer, (pH 7.4) at 37°C. 1, 3, 7, 14, 21, and 28 days after starting of the incubation, the amount of eluted doxorubicin hydrochloride was measured with a fluorescent spectrophotometer. The results are shown in Table 1, where the eluted amounts are represented by wt% to the amount of doxorubicin hydrochloride that is originally contained in the scleral plug (100%). The results were shown by the mean value of three test

samples.

Table 1

Day	The amount of eluted drug (%)
1	6.2
3	9.5
7	13.4
14	17.2
21	22.2
28	26.0

[0039] As shown in Table 1, doxorubicin hydrochloride was eluted slowly for four weeks. The results prove the sustained release effect of the drug, which is the object of the present invention. During the test period, disintegration of the plug was not observed.

(2) In vivo tests

[0040] Vitreous surgery was performed on one eye of each of ten pigmented rabbit. After two weeks, the scleral plug was inserted through the scleral incision and covered with conjunctiva. 1, 3, 5, 14, and 28 days after insertion, 0.2 ml aqueous humor was taken from vitreous cavity and stored at -80°C. The concentration of doxorubicin hydrochloride in the aqueous humor was measured by a high performance liquid chromatographic method. The results are shown in Table 2.

Table 2

Day	The concentration of drug (ng/ml), mean value of ten eyes
1	12.4
3	12.8
5	9.4
14	2.8
28	7.5

[0041] As shown in Table 2, doxorubicin hydrochloride was released gradually into the vitreous body for four weeks and the effective concentration was maintained (2 to 10 ng/ml) on each measuring day. Moreover, the concentration of the drug did not greatly rise at a time and accumulation of the drug in the vitreous body was not observed. The sustained release effect and the safety of the plug of the present invention were proved by this test.

(3) Toxicity tests

(Histopathologic change in tissues)

[0042] The scleral plug was inserted through a scleral incision into the vitreous body of a pigmented rabbit and covered with conjunctiva. After one or three months, an eyeball was excised to observe histopathologic change in eye tissues with a light microscope. No histopathologic change was observed and the safety of the plug was proved. After the excision of an eyeball, the condition of the plug was also checked to ascertain degradation of the scleral plug. The result proves that the scleral plug of the present invention is slowly degraded in a living body.

(Change in retinal functions)

[0043] Vitreous surgery was performed on one eye of each of two pigmented rabbits and then the scleral plug was inserted through the scleral incision and covered with conjunctiva. One month after insertion, an electroretinogram was recorded and compared with the electroretinogram recorded before vitreous surgery and also compared with the electroretinogram of the other eye, which was not treated by vitreous surgery. No change in b wave was observed in every case. The result proved that the scleral plug of the present invention did not affect the functions of retina.

INDUSTRIAL APPLICABILITY

[0044] The present invention provides a scleral plug that effectively plugs an incision after vitreous surgery and is not required to be removed. The scleral plug enables a drug to be administered safely and effectively in intraocular tissues, where drugs hardly distribute, by releasing a drug gradually for a long period. Moreover, it is possible to find another conventional methods for administering a drug, e.g., a method involving the use of a scleral incision which is formed during vitreous surgery.

Claims

1. Biodegradable medical device to deliver a medication into a vitreous body, which is made of lactic acid polymer and contains a drug, characterized in that the device is a nail-like scleral plug and has a head portion (1, 11) and a shaft portion (3, 13), wherein only the shaft portion (3, 13) of the scleral plug is insertable into the vitreous body through the sclera such that the head portion (1, 11) remains outside the vitreous body, whereby the shaft portion (3, 13) is absorbable into the vitreous body so that the drug is released gradually into the vitreous body from the scleral plug.
2. Biodegradable medical device according to claim 1, wherein the lactic acid polymer comprises lactic acid units and glycolic acid units.
3. Biodegradable medical device according to claim 2, wherein the weight-average molecular weight of the lactic acid copolymer is in the range from 10,000 to 1,000,000.
4. Biodegradable medical device according to claim 2 or 3, which contains a drug for use in promoting a recovery from damage after vitreous surgery, wherein said drug is released slowly into a vitreous body.
5. Biodegradable medical device according to any of claims 2 to 4, wherein the weight-average molecular weight of the copolymer is in the range from 10,000 to 100,000, and said plug is designed to release effective concentration of a drug for about one to two weeks.
6. Biodegradable medical device according to claim 5, wherein the weight-average molecular weight of the copolymer is in the range from 20,000 to 40,000.
7. Biodegradable medical device according to any of claims 2 to 4, wherein the weight-average molecular weight of the copolymer is in the range from 10,000 to 200,000, and said plug is designed to release effective concentration of a drug for about two weeks to one month.
8. Biodegradable medical device according to claim 7, wherein the weight-average molecular weight of the copolymer is in the range from 20,000 to 50,000.
9. Biodegradable medical device according to any of claims 2 to 4, wherein the weight-average molecular weight of the copolymer is in the range from 10,000 to 1,000,000, and said plug is designed to release effective concentration of a drug for one month or more.
10. Biodegradable medical device according to claim 9, wherein the weight-average molecular weight of the copolymer is in the range from 40,000 to 200,000.
11. Biodegradable medical device according to any of claims 2 to 10, wherein the molar ratio of lactic acid units and glycolic acid units is in the range from 50/50 to 80/20.
12. Biodegradable medical device according to claim 11, wherein the molar ratio of lactic units and glycolic acid units is in the range from 70/30 to 80/20.
13. Biodegradable medical device according to any

preceding claim which contains a drug for use in treating or preventing diseases of retina, wherein said drug is released gradually into a vitreous body.

14. Biodegradable medical device according to any preceding claim, wherein said lactic acid unit is in L, D, or DL form. 5
15. Biodegradable medical device according to claim 7 or 8, wherein said drug is an antitumor agent. 10
16. Biodegradable medical device according to claim 9 or 10, wherein said drug is an antiviral agent.
17. Biodegradable medical device according to claim 7 or 8, wherein said drug is an antiinflammatory agent. 15
18. Biodegradable medical device according to claim 3 or 6, wherein said drug is an antibiotics. 20
19. Biodegradable medical device according to claim 9 or 10, wherein said drug is an antifungal agent.

Patentansprüche

1. Biologisch abbaubare, medizinische Vorrichtung zur Verabreichung eines Medikamentes in einen Glaskörper, die aus einem Milchsäure-Polymer besteht und ein Arzneimittel enthält, dadurch gekennzeichnet, dass die Vorrichtung ein nagelähnlicher Sklerastopfen ist und einen Kopfabschnitt (1,11) und einen Schaftabschnitt (3,13) aufweist, wobei nur der Schaftabschnitt (3,13) des Sklerastopfens in den Glaskörper durch die Sklera einführbar ist, so dass der Kopfabschnitt (1,11) außerhalb des Glaskörpers verbleibt, wodurch der Schaftabschnitt (3,13) in dem Glaskörper absorbierbar ist, so dass das Arzneimittel allmählich aus dem Sklerastopfen in den Glaskörper freigesetzt wird. 30
2. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 1, wobei das Milchsäure-Polymer des Stopfens Milchsäure-Einheiten und Glycolsäure-Einheiten umfasst. 45
3. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 2, wobei die massegemittelte Molekülmasse des Milchsäure-Copolymers in dem Bereich von 10.000 bis 1.000.000 liegt. 50
4. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 2 oder 3, die ein Arzneimittel zur Verwendung bei der Förderung der Genesung von einem Schaden nach einer chirurgischen Glaskörperbehandlung enthält, wobei dieses Arzneimittel langsam in den Glaskörper freigesetzt wird. 55

5. Biologisch abbaubare, medizinische Vorrichtung nach einem der Ansprüche 2 bis 4, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 10.000 bis 100.000 liegt, und der Stopfen dafür bestimmt ist, eine effektive Konzentration eines Arzneimittels für etwa ein bis zwei Wochen freizusetzen.
6. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 5, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 20.000 bis 40.000 liegt.
7. Biologisch abbaubare, medizinische Vorrichtung nach einem der Ansprüche 2 bis 4, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 10.000 bis 200.000 liegt, und der Stopfen dafür bestimmt ist, eine effektive Konzentration eines Arzneimittels für etwa zwei Wochen bis einen Monat freizusetzen.
8. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 7, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 20.000 bis 50.000 liegt. 25
9. Biologisch abbaubare, medizinische Vorrichtung nach einem der Ansprüche 2 bis 4, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 10.000 bis 1.000.000 liegt, und der Stopfen dafür bestimmt ist, eine effektive Konzentration eines Arzneimittels für einen Monat oder mehr freizusetzen.
10. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 9, wobei die massegemittelte Molekülmasse des Copolymers in dem Bereich von 40.000 bis 200.000 liegt. 35
11. Biologisch abbaubare, medizinische Vorrichtung nach einem der Ansprüche 2 bis 10, wobei das molare Verhältnis der Milchsäure-Einheiten und der Glycolsäure-Einheiten in dem Bereich von 50/50 bis 80/20 liegt. 40
12. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 11, wobei das molare Verhältnis der Milchsäure-Einheiten und der Glycolsäure-Einheiten in dem Bereich von 70/30 bis 80/20 liegt.
13. Biologisch abbaubare, medizinische Vorrichtung nach einem der vorangehenden Ansprüche, die ein Arzneimittel zur Verwendung bei der Behandlung oder der Verhinderung von Krankheiten der Netzhaut enthält, wobei das Arzneimittel allmählich in den Glaskörper freigesetzt wird.
14. Biologisch abbaubare, medizinische Vorrichtung

nach einem der vorangehenden Ansprüche, wobei die Milchsäure in der L-, D- oder DL-Form vorliegt.

15. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 7 oder 8, wobei das Arzneimittel ein Antitumormittel ist. 5
16. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 9 oder 10, wobei das Arzneimittel ein antivirales Mittel ist. 10
17. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 7 oder 8, wobei das Arzneimittel ein entzündungshemmendes Mittel ist. 15
18. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 3 oder 6, wobei das Arzneimittel ein Antibiotikum ist.
19. Biologisch abbaubare, medizinische Vorrichtung nach Anspruch 9 oder 10, wobei das Arzneimittel ein antifungales Mittel ist. 20

Revendications

1. Dispositif médical biodégradable en vue de délivrer une médication dans un corps vitré, qui est fait d'un polymère d'acide lactique et contient un médicament, caractérisé en ce que le dispositif est un bouchon scléral semblable à un clou et comporte une partie de tête (1, 11) et une partie de tige (3, 13), dans lequel seule la partie de tige (3, 13) du bouchon scléral peut être introduite dans le corps vitré à travers la sclérotique de sorte que la partie de tête (1, 11) reste à l'extérieur du corps vitré, grâce à quoi la partie de tige (3, 13) peut être absorbée dans le corps vitré de sorte que le médicament soit libéré progressivement dans le corps vitré à partir du bouchon scléral. 25 30 35 40
2. Dispositif médical biodégradable selon la revendication 1, dans lequel le polymère d'acide lactique du bouchon comprend des unités d'acide lactique et des unités d'acide glycolique. 45
3. Dispositif médical biodégradable selon la revendication 2, dans lequel la masse moléculaire moyenne en poids du copolymère d'acide lactique se situe dans la plage allant de 10 000 à 1 000 000. 50
4. Dispositif médical biodégradable selon la revendication 2 ou 3, qui contient un médicament destiné à une utilisation pour favoriser une guérison d'un dommage après une chirurgie du vitré, dans lequel ledit médicament est libéré lentement dans un corps vitré. 55
5. Dispositif médical biodégradable selon l'une quel-

conque des revendications 2 à 4, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 10 000 à 100 000 et ledit bouchon est conçu pour libérer une concentration efficace d'un médicament pendant environ une à deux semaines.

6. Dispositif médical biodégradable selon la revendication 5, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 20 000 à 40 000.
7. Dispositif médical biodégradable selon l'une quelconque des revendications 2 à 4, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 10 000 à 200 000, et ledit bouchon est conçu afin de libérer une concentration efficace d'un médicament pendant environ deux semaines à un mois.
8. Dispositif médical biodégradable selon la revendication 7, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 20 000 à 50 000.
9. Dispositif médical biodégradable selon l'une quelconque des revendications 2 à 4, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 10 000 à 1 000 000, et ledit bouchon est conçu afin de libérer une concentration efficace d'un médicament pendant un mois ou plus.
10. Dispositif médical biodégradable selon la revendication 9, dans lequel la masse moléculaire moyenne en poids du copolymère se situe dans la plage allant de 40 000 à 200 000.
11. Dispositif médical biodégradable selon l'une quelconque des revendications 2 à 10, dans lequel le rapport molaire des unités d'acide lactique et des unités d'acide glycolique se situe dans la plage allant de 50/50 à 80/20.
12. Dispositif médical biodégradable selon la revendication 11, dans lequel le rapport molaire des unités d'acide lactique et des unités d'acide glycolique se situe dans la plage allant de 70/30 à 80/20.
13. Dispositif médical biodégradable selon l'une quelconque des revendications précédentes, qui contient un médicament destiné à une utilisation pour le traitement ou la prévention de lésions de la rétine, dans lequel ledit médicament est libéré progressivement dans un corps vitré.
14. Dispositif médical biodégradable selon l'une quelconque des revendications précédentes, dans

lequel ladite unité d'acide lactique est de forme L, D, ou DL.

15. Dispositif médical biodégradable selon la revendication 7 ou 8, dans lequel ledit médicament est un agent anti-tumoral. 5
16. Dispositif médical biodégradable selon la revendication 9 ou 10, dans lequel ledit médicament est un agent antiviral. 10
17. Dispositif médical biodégradable selon la revendication 7 ou 8, dans lequel ledit médicament est un agent anti-inflammatoire. 15
18. Dispositif médical biodégradable selon la revendication 3 ou 6, dans lequel ledit médicament est un antibiotique.
19. Dispositif médical biodégradable selon la revendication 9 ou 10, dans lequel ledit médicament est un agent antifongique. 20

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FIG. 1

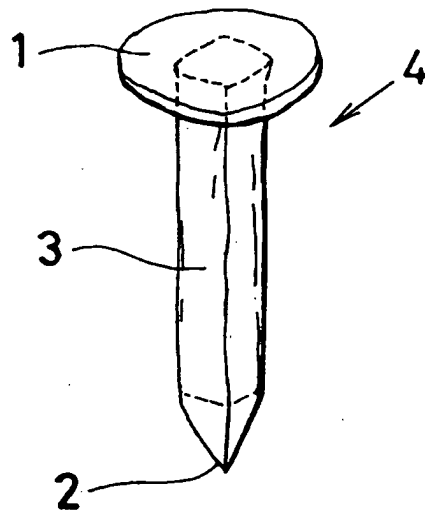


FIG. 2

